



# CALIFORNIA



## CALIFORNIA CANCER REGISTRY

Year Funded	Type of Program	Affiliation	Compliance with Public Law 102-515: Legislation and Regulations*	
1994	Enhancement	State Health Department	Authorizing legislation for a statewide registry: Yes	Regulations in place for data reporting, confidentiality, and use: All

\*Source: NPCR Legislative Database, January 2001.

### Cancer Burden

Cancer is a devastating disease. According to the American Cancer Society, in 2001 an estimated 553,400 Americans will die of cancer, and approximately 1.3 million new cancer cases will be diagnosed. These estimates do not include in situ (preinvasive) cancer of any site except urinary bladder or approximately 1 million cases of nonmelanoma skin cancer to be diagnosed this year. In California, an estimated 117,400 new cancer cases will be diagnosed, and approximately 51,200 will die of this disease in 2001. California ranks 41st highest overall in cancer mortality rates among the 50 states and Washington, D.C.\*

### The NPCR

Cancer surveillance involves the systematic collection, analysis, and use of cancer data. Information derived through surveillance is critical for directing effective cancer prevention and control programs. The Centers for Disease Control and Prevention's **National Program of Cancer Registries (NPCR)**, authorized by Congress in 1992 through the Cancer Registries Amendment Act (Public Law 102-515) and guided by the CDC, serves as the foundation of a national, comprehensive cancer control strategy. Statewide cancer registries provide critical data to help identify and monitor trends in cancer incidence and mortality over time; guide cancer control planning and

evaluation; help allocate health resources; and advance clinical, epidemiologic, and health services research.

CDC supports registries in 45 states, the District of Columbia, and three territories, representing 96% of the U.S. population.<sup>†</sup> Forty-five programs receive support to enhance existing registries. These states have on-going cancer data-collection activities, policies and procedures for central registry operations in place, and core staff employed. Four programs receive support to develop new registries; they are typically involved in laying the necessary groundwork for establishing a central, population-based registry.

With fiscal year 2001 appropriations of approximately \$36 million, CDC continues to support and enhance state cancer registries, and promotes appropriate uses of data. The CDC will also develop special research projects, such as studies of patterns of cancer care in specific populations and assessments of data for integration with geographic information systems.

### State Highlights

- The California Cancer Registry (CCR) was certified by the North American Association of Central Cancer Registries for completeness, timeliness, and quality of its 1997 data.
- The CCR is one of 11 registries participating in a special NPCR-

supported childhood cancer project to design, implement, and evaluate a method to use data from a state population-based central registry to compute expected numbers of incident cancer cases in children. The registry will evaluate completeness of its data and of other existing pediatric cancer databases, such as the Children's Oncology Group, by performing data linkage.

- California Teachers Study — This project is an on-going cohort study of more than 130,000 active and retired teachers to assess breast and other cancers, with an emphasis on modifiable risk factors.

### Contact Information

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\*Source: CDC's National Center for Health Statistics, vital statistics data, underlying cause of death, 1993-1997.

<sup>†</sup>A map of the participating NPCR programs can be found at <http://www.cdc.gov/cancer/npcr/statecon.htm>.